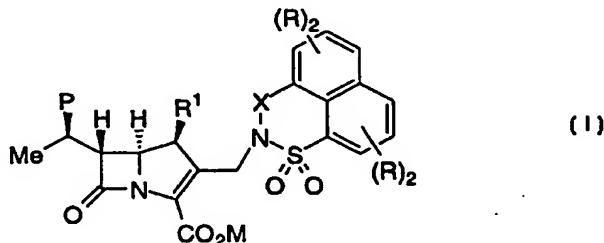




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C07D 477/14		A1	(11) International Publication Number: WO 99/67242 (43) International Publication Date: 29 December 1999 (29.12.99)
<p>(21) International Application Number: PCT/US99/14235</p> <p>(22) International Filing Date: 23 June 1999 (23.06.99)</p> <p>(30) Priority Data: 60/090,613 25 June 1998 (25.06.98) US</p> <p>(71) Applicant (for all designated States except US): MERCK & CO., INC. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).</p> <p>(72) Inventors; and</p> <p>(75) Inventors/Applicants (for US only): RATCLIFFE, Ronald, W. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). DYKSTRA, Kevin, D. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). BLIZZARD, Timothy, A. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).</p> <p>(74) Common Representative: MERCK & CO., INC.; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).</p>		<p>(81) Designated States: AE, AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ZA, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>	

(54) Title: NAPHTHO[1,8-DE]THIASIN-2-YL METHYL CARBAPENEM ANTIBACTERIALS



(57) Abstract

Compounds of formula (I) as well as pharmaceutically acceptable salts thereof are disclosed.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon	KR	Republic of Korea	PL	Poland		
CN	China	KZ	Kazakhstan	PT	Portugal		
CU	Cuba	LC	Saint Lucia	RO	Romania		
CZ	Czech Republic	LI	Liechtenstein	RU	Russian Federation		
DE	Germany	LK	Sri Lanka	SD	Sudan		
DK	Denmark	LR	Liberia	SE	Sweden		
EE	Estonia			SG	Singapore		

**NAPHTHO[1,8-DE]THIASIN-2-YL METHYL CARBAPENEM
ANTIBACTERIALS**

5

BACKGROUND OF THE INVENTION

The present invention relates to carbapenem antibacterial agents in which the carbapenem nucleus is substituted at the 2-position with a 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl group or

10 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl group linked through a CH₂ group.

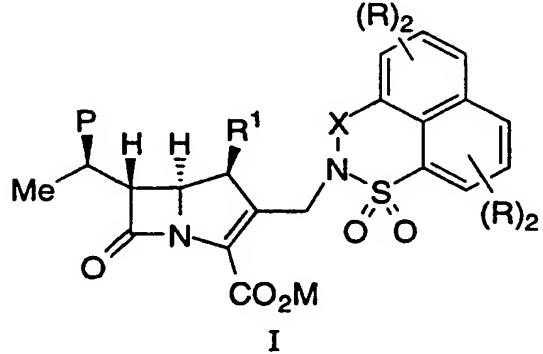
The carbapenems of the present invention are useful against gram positive microorganisms, especially methicillin resistant *Staphylococcus aureus* (MRSA), methicillin resistant *Staphylococcus epidermidis* (MRSE), and methicillin resistant coagulase negative *Staphylococci* (MRCNS). The antibacterial compounds of the present invention thus comprise an important contribution to therapy for treating infections caused by these difficult to control pathogens. There is an increasing need for agents effective against such pathogens

20 (MRSA/MRCNS) which are at the same time relatively free from undesirable side effects.

SUMMARY OF THE INVENTION

The compounds of the invention are represented by formula

25 I:



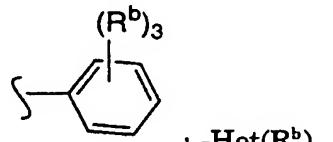
including pharmaceutically acceptable salts thereof, wherein:

R^1 represents H or methyl;

CO_2M represents a carboxylic acid, a pharmaceutically acceptable carboxylic acid salt, carboxylate anion, a pharmaceutically acceptable ester group or a carboxylic acid protected by a protecting group;

10 X represents CH_2 or $C=O$;

P represents hydrogen, hydroxyl, F or hydroxyl protected by a hydroxyl-protecting group;

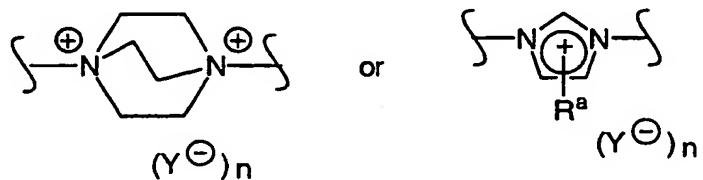


Each R independently represents R^b , $-Het(R^b)_3$,

15 C_{2-6} alkenyl, or a group $L-Q-R^a$ with the proviso that only one R group of the type $L-Q-R^a$ can be present or one R group may be taken with L , if present, and any intervening atoms to represent a 5-6 membered ring;

20 L is C_{1-4} straight or branched alkylene, uninterrupted, interrupted or terminated by 1-2 of O, S, NR^a , $C(O)$, CO_2 and $C(O)NR^a$;

Q represents:



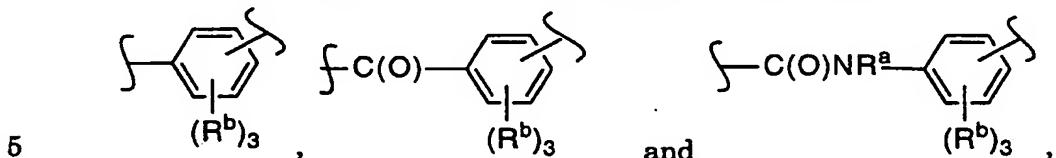
25

Y^- is a charge balancing group;

n is a value from 0 to 2 selected to maintain overall charge neutrality;

R^a is H or C₁₋₆ alkyl;

R^q is C₁₋₆ alkyl, straight or branched, uninterrupted, interrupted or terminated by 1-2 of O, S, NR^a, C(O), C(O)O, C(O)NR^a, -CH=CH-, -Het(R^b)₃-, -C(O)Het(R^b)₃-, -C(O)NR^aHet(R^b)₃-,

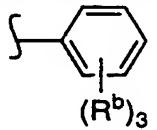


said R^q being unsubstituted or substituted with 1-3 R^c groups;

Het is a heteroaryl group;

10 each R^b is independently selected from H, halo, OR^a, OC(O)R^a, C(O)R^a, CN, C(O)NR^aR^d, NO₂, NR^aR^d, SO₂NR^aR^d and C₁₋₄ alkyl unsubstituted or substituted with 1-3 groups selected from R^e;

15 each R^c is independently selected from halo, OR^f, OC(O)R^f, SR^f, S(O)R^f, SO₂R^f, CN, C(O)R^f, CO₂R^f, NR^fR^g, N⁺R^aR^fR^gZ⁻, C(O)NR^aR^f, -Het(R^b)₃, C(=N⁺R^aR^f)R^gZ⁻, C(=N⁺R^aR^f)NR^aR^fZ⁻, NR^aC(=N⁺R^aR^f)R^gZ⁻, NR^aC(=N⁺R^aR^f)NR^aR^fZ⁻, heteroarylium(R^b)₃Z⁻, SO₂NR^aR^f, OC(O)NR^aR^f, NR^aC(O)R^f, NR^aC(O)NR^aR^f, and



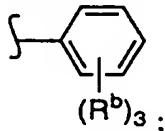
or in the alternative, when 2 or more R^c groups are present, 2 R^c groups may be taken together with any intervening atoms to form a 3-6 membered carbocyclic ring, optionally interrupted with 1-3 of O, S, NR^g, and C(O), said ring being unsubstituted or substituted with 1-3 R^e groups;

R^d is H or C₁₋₄ alkyl, or R^a and R^d taken together with any intervening atoms represent a 4-6 membered ring;

R^e is selected from halo, OR^a , NR^aR^d and $CONR^aR^d$;

R^f is H; C1-6 straight or branched chain alkyl, unsubstituted or substituted with 1-3 R^e groups; - $Het(R^b)3$; C3-6

5 cycloalkyl, unsubstituted or substituted with 1-3 R^e groups, and



or R^a and R^f together with an intervening atoms form a 4-6 membered ring, optionally interrupted by O, S, NR^a or $C(O)$; and

10 R^g is H, C1-6 alkyl, unsubstituted or substituted with 1-3 R^e groups; C3-6 cycloalkyl, unsubstituted or substituted with 1-3 R^e groups; $C(=N^+R^aR^f)R^aZ^-$ or $C(=N^+R^aR^f)NR^aR^fZ^-$;

15 or R^f and R^g together with any intervening atoms form a 4-6 membered ring optionally interrupted by O, S, NR^a or $C(O)$; and

20 Z^- is a charge balancing group selected from Y^- , a monovalent anion such as acetate, benzoate, bromide, chloride or the like, or an internal anion such as CO_2M , where M is a negative charge.

DETAILED DESCRIPTION OF THE INVENTION

The invention is described herein in detail using the terms defined below unless otherwise specified.

25 Carboxylate anion refers to a negatively charged group $-COO^-$.

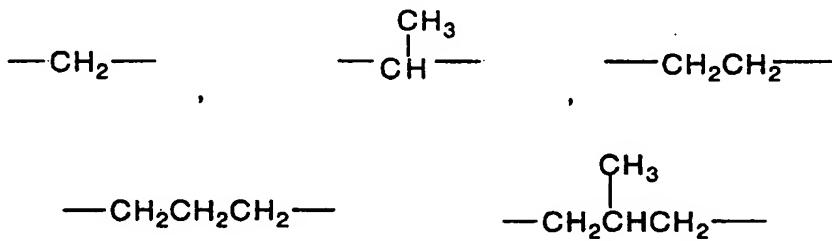
The term "alkyl" refers to a monovalent alkane (hydrocarbon) derived radical containing from 1 to 15 carbon atoms unless otherwise defined. It may be straight or branched.

30 Preferred alkyl groups include methyl, ethyl, propyl, isopropyl, butyl and t-butyl. When substituted, alkyl groups may be substituted with up to 3 substituent groups, selected from R^c or R^e as defined, at any available

point of attachment. When the alkyl group is said to be substituted with an alkyl group, this is used interchangeably with "branched alkyl group".

5 Cycloalkyl is a species of alkyl containing from 3 to 15 carbon atoms, without alternating or resonating double bonds between carbon atoms. It may contain from 1 to 4 rings which are fused. Preferred cycloalkyl groups are cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl. When substituted, cycloalkyl groups may be substituted with up to 3 substituents selected from R^c or R^e as defined.

10 A C_{1-4} alkylene group refers to an alkyl group which is attached through two bonds to two different atoms or substituents. The two bonds on the alkylene group can be on the same carbon atom or on different carbon atoms. See, e.g., the following:



15

The term "alkenyl" refers to a hydrocarbon radical straight, branched or cyclic containing from 2 to 10 carbon atoms and at least one carbon to carbon double bond. Preferred alkenyl groups include ethenyl, 20 propenyl, butenyl and cyclohexenyl.

The term "alkynyl" refers to a hydrocarbon radical straight or branched, containing from 2 to 10 carbon atoms and at least one carbon to carbon triple bond. Preferred alkynyl groups include ethynyl, propynyl and butynyl.

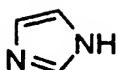
25 Aryl refers to aromatic rings e.g., phenyl, substituted phenyl and the like, as well as rings which are fused, e.g., naphthyl, phenanthrenyl and the like. An aryl group thus contains at least one ring having at least 6 atoms, with up to five such rings being present, containing up to 22 atoms therein, with alternating (resonating) double 30 bonds between adjacent carbon atoms. The preferred aryl groups are

phenyl, naphthyl and phenanthrenyl. Aryl groups may likewise be substituted as defined. Preferred substituted aryls include phenyl and naphthyl.

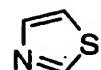
The term "heteroaryl" (Het) refers to a monocyclic aromatic hydrocarbon group having 5 or 6 ring atoms, or a bicyclic aromatic group having 8 to 10 atoms, containing at least one heteroatom, O, S or N, in which a carbon or nitrogen atom is the point of attachment, and in which one or two additional carbon atoms is optionally replaced by a heteroatom selected from O or S, and in which from 1 to 3 additional carbon atoms are optionally replaced by nitrogen heteroatoms, said heteroaryl group being optionally substituted as described herein. Examples of this type are pyrrole, pyridine, oxazole, thiazole and oxazine. Additional nitrogen atoms may be present together with the first nitrogen and oxygen or sulfur, giving, e.g., thiadiazole. Examples include the following:



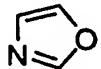
pyrrole (pyrrolyl)



imidazole (imidazolyl)



thiazole (thiazolyl)



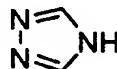
oxazole (oxazolyl)



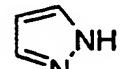
furan (furyl)



thiophene (thienyl)



triazole (triazolyl)



pyrazole (pyrazolyl)



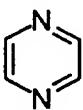
isoxazole (isoxazolyl)



isothiazole (isothiazolyl)



pyridine (pyridinyl)



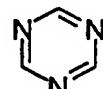
pyrazine (pyrazinyl)



pyridazine (pyridazinyl)



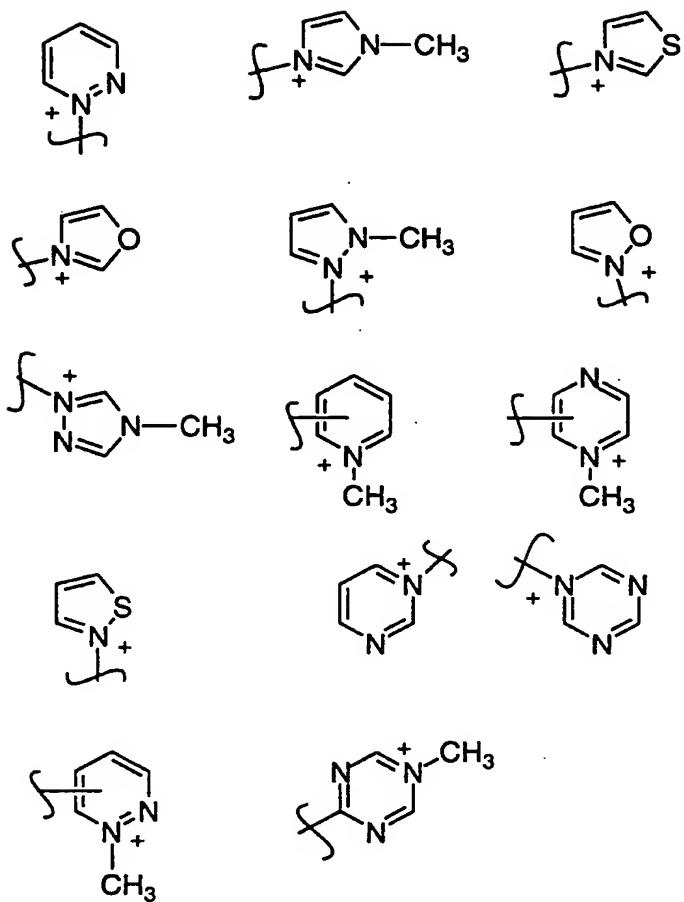
pyrimidine (pyrimidinyl)



triazine (triazinyl)

The group L-Q-Rq, if present, is attached to either of the two phenyl rings of the 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl group,
 5 provided that no more than one L-Q-Rq group is present.

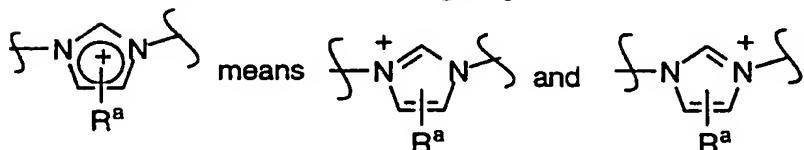
Heteroarylium refers to heteroaryl groups bearing a quaternary nitrogen atom and thus a positive charge. Examples include the following:



When a charge is shown on a particular nitrogen atom in a ring which contains one or more additional nitrogen atoms, it is
 5 understood that the charge may reside on a different nitrogen atom in the ring by virtue of charge resonance that occurs.



The imidazolium group:



The term "heterocycloalkyl" refers to a cycloalkyl group

5 (nonaromatic) in which one of the carbon atoms in the ring is replaced by a heteroatom selected from O, S or N, and in which up to three additional carbon atoms may be replaced by hetero atoms.

10 The terms "quaternary nitrogen" and "positive charge" refer to tetravalent, positively charged nitrogen atoms including, e.g., the positively charged nitrogen in a tetraalkylammonium group (e. g. tetramethylammonium), heteroarylium, (e.g., N-methyl-pyridinium), basic nitrogens which are protonated at physiological pH, and the like. Cationic groups thus encompass positively charged nitrogen-containing groups, as well as basic nitrogens which are protonated at physiologic pH.

15 The term "heteroatom" means O, S or N, selected on an independent basis.

Halogen and "halo" refer to bromine, chlorine, fluorine and iodine.

20 When a group is termed "substituted", unless otherwise indicated, this means that the group contains from 1 to 4 substituents thereon.

25 When a functional group is termed "protected", this means that the group is in modified form to preclude undesired side reactions at the protected site. Suitable protecting groups for the compounds of the

present invention will be recognized from the present application taking into account the level of skill in the art, and with reference to standard textbooks, such as Greene, T. W. et al. Protective Groups in Organic Synthesis Wiley, New York (1991). Examples of suitable protecting groups are contained throughout the specification.

In some of the carbapenem compounds of the present invention, M is a readily removable carboxyl protecting group, and/or P represents a hydroxyl which is protected by a hydroxyl-protecting group. Such conventional protecting groups consist of groups which are used to 10 protectively block the hydroxyl or carboxyl group during the synthesis procedures described herein. These conventional blocking groups are readily removable, i.e., they can be removed, if desired, by procedures which will not cause cleavage or other disruption of the remaining portions of the molecule. Such procedures include chemical and 15 enzymatic hydrolysis, treatment with chemical reducing or oxidizing agents under mild conditions, treatment with a transition metal catalyst and a nucleophile and catalytic hydrogenation.

Examples of carboxyl protecting groups include allyl, benzhydryl, 2-naphthylmethyl, benzyl, silyl such as 20 t-butyldimethylsilyl (TBDMS), phenacyl, p-methoxybenzyl, o-nitrobenzyl, p-methoxyphenyl, p-nitrobenzyl, 4-pyridylmethyl and t-butyl.

Examples of suitable hydroxy protecting groups include 25 triethylsilyl, t-butyldimethylsilyl, o-nitrobenzyloxycarbonyl, p-nitrobenzyloxycarbonyl, benzyloxycarbonyl, allyloxycarbonyl, t-butyloxycarbonyl, 2,2,2-trichloroethyloxycarbonyl and the like.

The carbapenem compounds of the present invention are useful per se and in their pharmaceutically acceptable salt and ester forms for the treatment of bacterial infections in animal and human 30 subjects. The term "pharmaceutically acceptable ester, salt or hydrate," refers to those salts, esters and hydrated forms of the compounds of the present invention which would be apparent to the pharmaceutical chemist. i.e., those which are substantially non-toxic and which may favorably affect the pharmacokinetic properties of said compounds, such 35 as palatability, absorption, distribution, metabolism and excretion.

Other factors, more practical in nature, which are also important in the selection, are cost of the raw materials, ease of crystallization, yield, stability, solubility, hygroscopicity and flowability of the resulting bulk drug. Conveniently, pharmaceutical compositions may be prepared

5 from the active ingredients in combination with pharmaceutically acceptable carriers. Thus, the present invention is also concerned with pharmaceutical compositions and methods of treating bacterial infections utilizing as an active ingredient the novel carbapenem compounds.

10 With respect to $-\text{CO}_2\text{M}$, which is attached to the carbapenem nucleus at position 3, this represents a carboxylic acid group (M represents H), a carboxylate anion (M represents a negative charge), a pharmaceutically acceptable carboxylic acid salt (M represents a salt forming group), a pharmaceutically acceptable ester (M represents an ester forming group) or a carboxylic acid protected by a protecting group (M represents a carboxyl protecting group).

15

The pharmaceutically acceptable salts referred to above may take the form $-\text{COOM}$, where M is a negative charge, which is balanced by a counterion, e.g., an alkali metal cation such as sodium or potassium. Other pharmaceutically acceptable counterions may be calcium, magnesium, zinc, ammonium, or alkylammonium cations such as tetramethylammonium, tetrabutylammonium, choline, triethylhydroammonium, meglumine, triethanolhydroammonium, etc.

20 The pharmaceutically acceptable salts referred to above also include acid addition salts. Thus, the Formula I compounds can be used in the form of salts derived from inorganic or organic acids. Included among such salts are the following: acetate, adipate, alginate, aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, glucoheptanoate, glycerophosphate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate,

pamoate, pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, tosylate and undecanoate.

Acid addition salts of the compounds of formula I include compounds that contain a protonated, basic moiety in R. Compounds 5 containing a basic moiety in R are capable of protonation in aqueous media near pH 7, so that the basic moiety can exist as an equilibrium mixture of its neutral form and acid addition (protonated) form. The more basic the group, the greater the degree of protonation near pH 7. For example, $-\text{NR}^f\text{RG}$ would likely be present in its protonated form 10 $(-\text{N}^+\text{HR}^f\text{RGZ}^-)$ at the appropriate pH, where Z^- is a charge balancing group. All such compounds are included in the present invention.

The pharmaceutically acceptable esters are such as would be readily apparent to a medicinal chemist, and include, for example, those described in detail in U.S. Pat. No. 4,309,438. Included within 15 such pharmaceutically acceptable esters are those which are hydrolyzed under physiological conditions, such as pivaloyloxymethyl, acetoxyethyl, phthalidyl, indanyl and methoxymethyl, and others described in detail in U.S. Pat. No. 4,479,947. These are also referred to as "biolabile esters".

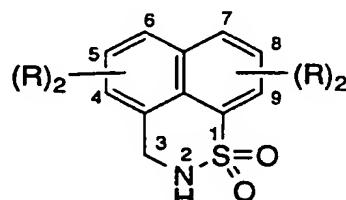
20 Biolabile esters are biologically hydrolizable, and may be suitable for oral administration, due to good absorption through the stomach or intestinal mucosa, resistance to gastric acid degradation and other factors. Examples of biolabile esters include compounds in which M represents an alkoxyalkyl, alkylcarbonyloxyalkyl, 25 alkoxy carbonyloxyalkyl, cycloalkoxyalkyl, alkenyloxyalkyl, aryloxyalkyl, alkoxyaryl, alkylthioalkyl, cycloalkylthioalkyl, alkenylthioalkyl, arylthioalkyl or alkylthioaryl group. These groups can be substituted in the alkyl or aryl portions thereof with acyl or halo groups. The following M species are examples of biolabile ester forming moieties.: 30 acetoxyethyl, 1-acetoxyethyl, 1-acetoxypropyl, pivaloyloxymethyl, 1-isopropylcarbonyloxyethyl, 1-cyclohexylcarbonyloxyethyl, phthalidyl and (2-oxo-5-methyl-1,3-dioxolen-4-yl)methyl.

35 Z^- and Y^- can be present or absent as necessary to maintain the appropriate charge balance. When present, these represent pharmaceutically acceptable counterions. Most anions derived from

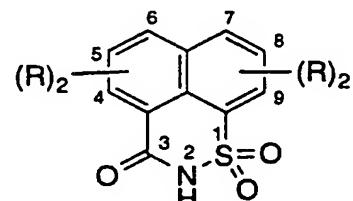
inorganic or organic acids are suitable. Representative examples of such counterions are the following: acetate, adipate, aminosalicylate, anhydromethylenecitrate, ascorbate, aspartate, benzoate, benzenesulfonate, bromide, citrate, camphorate, camphorsulfonate, 5 chloride, estolate, ethanesulfonate, fumarate, glucoheptanoate, gluconate, glutamate, lactobionate, malate, maleate, mandelate, methanesulfonate, pantothenate, pectinate, phosphate/diphosphate, polygalacturonate, propionate, salicylate, stearate, succinate, sulfate, tartrate and tosylate. Other suitable anionic species will be apparent to 10 the ordinarily skilled chemist.

Likewise, when more than one negative charge is necessary to maintain charge neutrality, the counterion indicator may represent a species with more than one negative charge, such as malonate, tartrate or ethylenediaminetetraacetate (EDTA), or two or more monovalent 15 anions, such as chloride, etc. When a multivalent negatively charged counterion is present with a carbapenem which bears a net single positive charge, an appropriate number of carbapenem molecules can be found in association therewith to maintain the overall charge balance and neutrality.

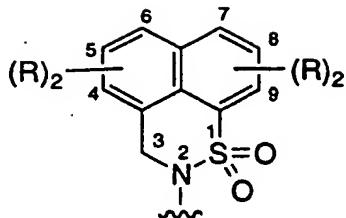
20 Numbering and nomenclature used in naming the sultam side chains are as follows:



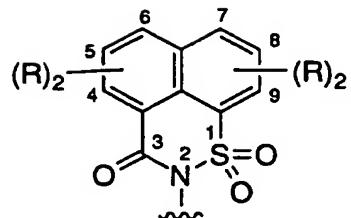
2,3-dihydro-naphtho[1,8-de][1,2]thiazine
1,1-dioxide



1,1-dioxo-2,3-dihydro-
naphtho[1,8-de][1,2]thiazin-3-one



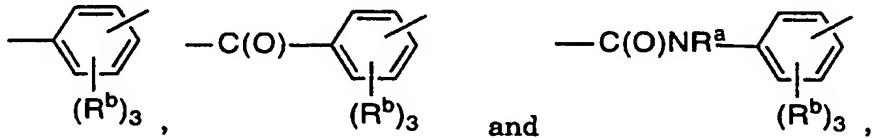
1,1-dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl



1,1,3-trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl

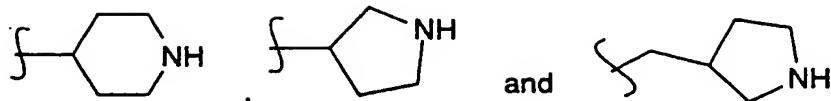
When L is a C₁₋₄ straight or branched alkylene group that is interrupted or terminated by 1-2 of O, S, NR^a, C(O), CO₂ and C(O)NR^a, 5 the interrupting/terminating moiety or moieties can be at either end of the alkylene group, as well as interrupting the alkylene group when 2-4 carbon atoms are present. When 2 such groups are present, they may be separate or together. Hence, interrupting or terminating groups such as OC(O) and OCO₂ are included.

10 Similarly, when R^q is C₁₋₆ alkyl, straight or branched, interrupted or terminated by 1-2 of O, S, NR^a, C(O), C(O)O, C(O)NR^a, -CH=CH-, -Het(R^b)₃-, -C(O)Het(R^b)₃-, -C(O)NR^aHet(R^b)₃-,



15 said R^q being unsubstituted or substituted with 1-3 R^c groups, the interrupting/terminating groups may be separate or together, and may be at the end or ends of the alkyl group, and further may be between the alkyl group and a substituent R^c.

20 When an R^q is substituted with at least 2 R^c groups, these may be taken in combination with any intervening atoms to represent a 3-6 membered carbocyclic ring, said ring being optionally interrupted by 1-3 of O, S, NR^c and C(O), and unsubstituted or substituted with 1-3 R^c groups. Examples of groups which are represented by two R^c groups in combination include the following:

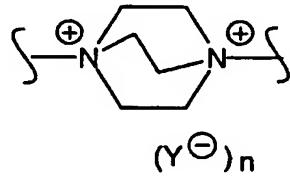


A subset of compounds of formula I which is of interest relates to those compounds where R¹ represents methyl. Within this subset, all other variables are as originally defined.

5 Another subset of compounds of formula I which is of interest relates to those compounds where CO₂M represents a carboxylate anion. Hence, M in this instance represents a negative charge which is balanced by a positively charged group, such as in the positively charged Q group. Likewise, if the positively charged Q group 10 contains more than one positive charge, a negatively charged counterion may be present which in combination with the carboxylate anion, provides overall charge neutrality.

Another subset of compounds of formula I that is of interest relates to those compounds where P represents hydroxyl or hydroxyl 15 protected by a hydroxyl protecting group. Within this subset, all other variables are as originally defined.

Another subset of compounds of formula I that is of interest relates to compounds where one R represents L-Q-R_q and all other R groups represent H, halo or C₁₋₄ alkyl unsubstituted or substituted with 20 1-3 groups selected from R^c. Within this subset, all other variables are as originally defined. An embodiment of this subset is realized when L represents -CH₂- or -CH₂CH₂- . Another embodiment of this subset is realized when Q represents



25 wherein Y⁻ represents a charge balancing group, and n is 0 to 2. Within this subset, all other variables are as originally defined. Still another embodiment of this subset is realized when R_q is straight or branched C₁₋₆ alkyl, substituted with 1-3 R^c groups. Within this subset, all other variables are as originally defined.

Another subset of compounds of formula I that is of interest relates to compounds where X is CH₂. Within this subset, all other variables are as originally defined.

5 Another subset of compounds of formula I that is of interest relates to compounds where X is CO. Within this subset, all other variables are as originally defined.

10 Another subset of compounds of formula I that is of interest relates to compounds where R is H, halo or C₁₋₄ alkyl unsubstituted or substituted with 1-3 groups selected from Re. Within this subset, all other variables are as originally defined.

A preferred subset of compounds of formula I which is of interest relates to those compounds wherein:

R¹ represents CH₃;

CO₂M represents a carboxylate anion;

15 P represents hydroxyl or hydroxyl protected by a hydroxyl protecting group; and

one R is L-Q-R^q and each remaining R is independently H, halo or C₁₋₄ alkyl unsubstituted or substituted with 1-3 groups selected from Re; all other variables are as originally described.

20 In one embodiment of this invention

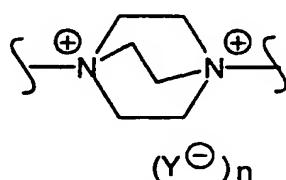
R^a is H or C₁₋₆ alkyl;

R^d is H or C₁₋₄ alkyl, or R^a and R^d taken together with any intervening atoms represent a 4-6 membered ring;

R^e is halo, OR^a, NR^aR^d or CONR^aR^d;

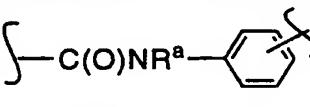
25 L represents -CH₂- or -CH₂CH₂-;

Q represents



wherein Y⁻ represents a charge balancing group, and n is 0 to 2; and

R^q is straight or branched C₁₋₆ alkyl, optionally interrupted

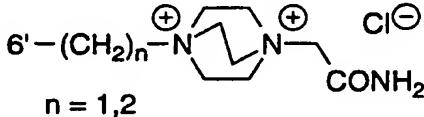
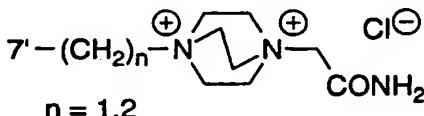
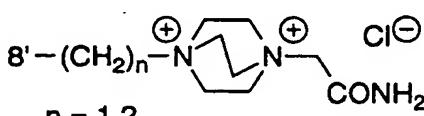
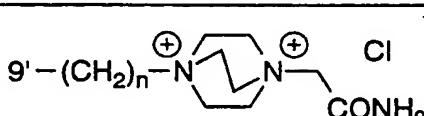
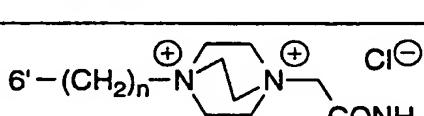
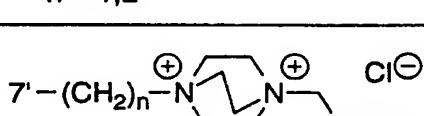
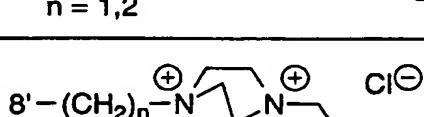
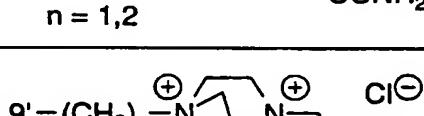
by C(O)NR^a or  and
substituted with 1-3 R^c groups, and
R^c is as originally defined.

5

Representative examples of compounds of the invention are found in Table I.

Table I

$P = H, F$ or OH , and $R^1 = Methyl$ or H			
	X	Substituent(s) R	M
1	CO	H	Na
2	CH ₂	H	Na
3	CO	7'-(CH ₂) _n -OH	Na
4	CO	7'-(CH ₂) _n - ⁺ n = 1,2	\ominus
5	CO	8'-(CH ₂) _n - ⁺ n = 1,2	\ominus
6	CH ₂	7'-(CH ₂) _n - ⁺ n = 1,2	\ominus
7	CO	7'-(CH ₂) _n - ⁺ -CH ₂ CH ₂ OH n = 1,2	\ominus

	X	Substituent(s) R	M
8	CO	 n = 1,2	⊖
9	CO	 n = 1,2	⊖
10	CO	 n = 1,2	⊖
11	CO	 n = 1,2	⊖
12	CH ₂	 n = 1,2	⊖
13	CH ₂	 n = 1,2	⊖
14	CH ₂	 n = 1,2	⊖
15	CH ₂	 n = 1,2	⊖

⊖

	X	Substituent(s) R	M
16	CO	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2)_3\text{OH} \text{ Cl}^-$ $n = 1,2$	\ominus
17	CO	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2)_3\text{OH} \text{ Cl}^-$ $n = 1,2$	\ominus
18	CH ₂	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2)_3\text{OH} \text{ Cl}^-$ $n = 1,2$	\ominus
19	CO	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2)_3\text{NH}_3^+ \text{ 2 Cl}^-$ $n = 1,2$	\ominus
20	CO	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2)_3\text{NH}_3^+ \text{ 2 Cl}^-$ $n = 1,2$	\ominus
21	CH ₂	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2)_3\text{NH}_3^+ \text{ 2 Cl}^-$ $n = 1,2$	\ominus
22	CO	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2\text{C}(=\text{O})\text{NH-CH}_2\text{CH}_2\text{NH}_3^+ \text{ 2 Cl}^-$ $n = 1,2$	\ominus
23	CO	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2)_2\text{N}^+(\text{CH}_2\text{C}(=\text{O})\text{NH-CH}_2\text{CH}_2\text{NH}_3^+ \text{ 2 Cl}^-$ $n = 1,2$	\ominus

	X	Substituent(s) R	M
24	CH ₂	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+(\text{CH}_2\text{CNH-CH}_2\text{CH}_2\text{NH}_3)_2\text{Cl}^-$ $n = 1,2$	⊖
25	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+(\text{CH}_2\text{CNH-CH}_2\text{CH}_2\text{NH}_3)_2\text{Cl}^-$ $n = 1,2$	⊖
26	CO	$6'\text{-Br, } 8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+\text{CONH}_2\text{Cl}^-$ $n = 1,2$	⊖
27	CO	$6'\text{-Br, } 8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+\text{CONH}_2\text{Cl}^-$ $n = 1,2$	⊖
28	CH ₂	$6'\text{-Br, } 8'-(\text{CH}_2)_n\text{N}^+(\text{CH}_2)_2\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+\text{CONH}_2\text{Cl}^-$ $n = 1,2$	⊖
29	CO	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+(\text{CH}_2\text{CONH}_2)_2\text{Cl}^-$ $n = 1,2$	⊖
30	CH ₂	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+(\text{CH}_2\text{CONH}_2)_2\text{Cl}^-$ $n = 1,2$	⊖
31	CO	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4)_2\text{N}^+(\text{CH}_2\text{CONH}_2)_2\text{Cl}^-$ $n = 1,2$	⊖

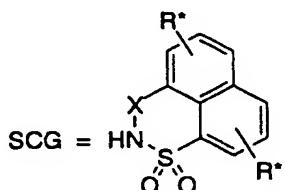
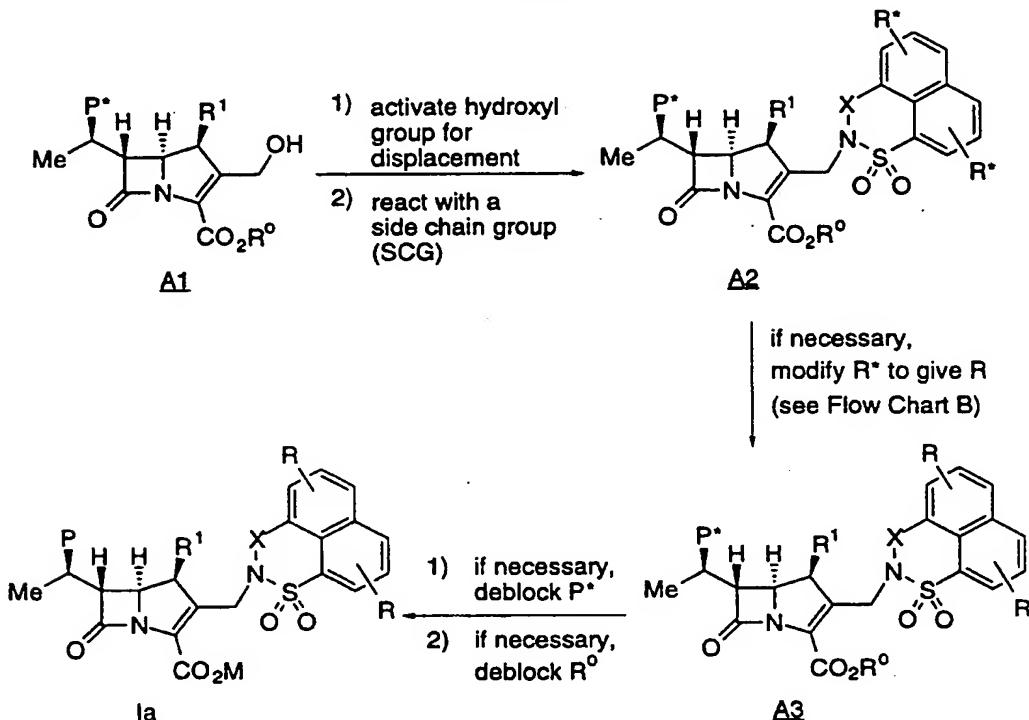
	X	Substituent(s) R	M
32	CH ₂	$7'-(\text{CH}_2)_n\text{N}^+ \text{---} \text{CH}_2\text{CH}_2\text{OH}$ $n = 1,2$	Cl^-
33	CO	$8'-(\text{CH}_2)_n\text{N}^+ \text{---} \text{CH}_2\text{CH}_2\text{OH}$ $n = 1,2$	Cl^-
34	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+ \text{---} \text{CH}_2\text{CH}_2\text{OH}$ $n = 1,2$	Cl^-
35	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+ \text{---} \text{CH}_3$ $n = 1,2$	Cl^-
36	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+ \text{---} \text{C}_2\text{H}_4\text{OH}$ $n = 1,2$	Cl^-
37	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+ \text{---} \text{C}_2\text{H}_4\text{NH}_3^+$ $n = 1,2$	2Cl^-

The compounds of the present invention are prepared by

5 reacting a suitably protected, activated 2-hydroxymethyl-carbapen-2-em-3-carboxylate with a 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl, modifying the thus-introduced side chain as desired, and then removing any protecting groups which are present to afford the desired final product.

The process is illustrated using the general conditions shown in the accompanying flow charts.

Flow Chart A

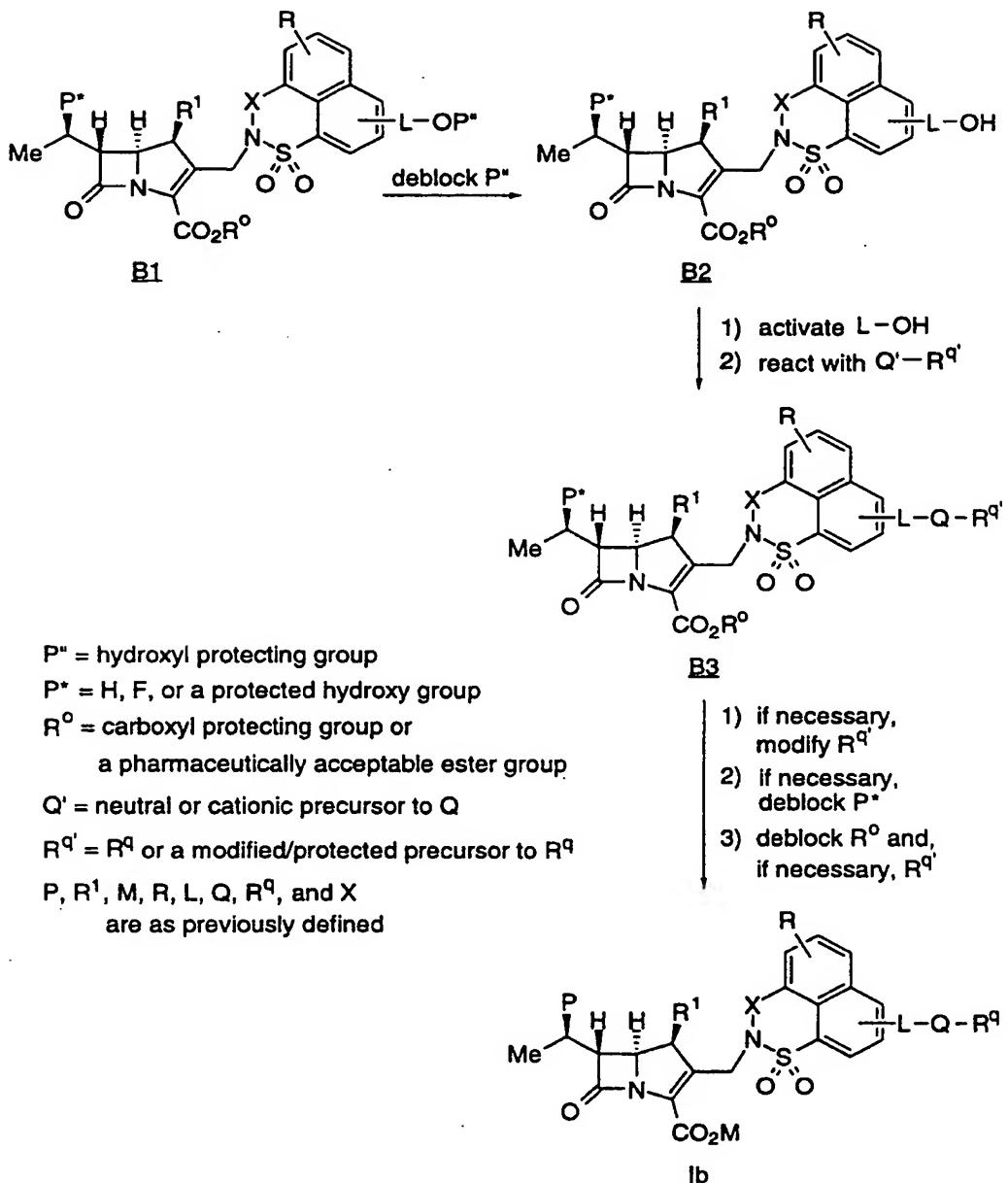


P^* = H, F, or a protected hydroxy group

R^o = carboxyl protecting group or a pharmaceutically acceptable ester group

R^* = R or a precursor to R

P , R^1 , M , R , and X are as previously defined

Flow Chart B

With reference to Flow Charts A and B above, $\text{P}, \text{R}^1, \text{R}, \text{M}, \text{X}, \text{L}, \text{Q}$ and R^{q} are as defined with respect to the compounds of formula I.

R^0 represents a carboxyl protecting group.

R^* represents a group selected from R or a group that is modified as necessary in the course of the synthesis of a compound of formula I to afford a member of R , thus R^* can be viewed as R or a precursor to R .

P^* represents a hydroxyl protecting group.

$Q'-R^q$ represents a neutral or monocationic group that reacts with the intermediate $B2$ (upon activation of $B2$) in a manner which results in the incorporation in the final product of the member of the group defined as $Q-R^q$ above, thus $Q'-R^q$ may be viewed as a precursor for $Q-R^q$.

R^q represents a group selected from R^q or a group that is modified or deprotected as necessary in the course of the synthesis of a compound of formula I so as to afford a member of R^q , thus R^q can be viewed as R^q or as a precursor to R^q .

The 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl side chain group (SCG) used in the synthesis of the compounds of the present invention have, in some cases, been described in the chemical literature. In other cases, precursor compounds which may be readily converted to the requisite SCG have been described in the literature. In cases where the requisite SCG is not known in the literature it is necessary to synthesize the SCG by a newly developed synthesis. One skilled in the art can adapt a previously published synthesis of an analogous SCG to prepare the requisite compound in a straightforward manner without undue experimentation.

The 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl side chain group (SCG) is initially reacted with a suitably protected carbapen-2-em-3-carboxylate having an activated hydroxymethyl group at the 2-position. The carbapenem nucleus having a $-CH_2OH$ substituent at position 2 can be obtained in accordance with Schmitt, S. M. et al., *J. Antibiotics* 41(6): 780-787 (1988), the teachings of which are incorporated herein by reference.

The carboxylic acid group at C-3 of the carbapenem is generally protected as a carboxyl protecting group such as p-nitrobenzyl (PNB), allyl, p-methoxybenzyl, trichloroethyl, 2-trimethylsilylethyl, and the like. Furthermore, the hydroxyl group of the 6-(hydroxyethyl) side-chain is optionally protected with a hydroxyl protecting group such as trimethylsilyl (TMS), triethylsilyl (TES), tert-butyldimethylsilyl (TBDMS), tert-butyldiphenylsilyl (TBDPS), acetyl, allyloxycarbonyl, 2-trimethylsilylethoxy carbonyl, 2-trichloroethoxycarbonyl and the like.

The addition of 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl side chain group (SCG) to the carbapenem is accomplished by treating a solution of the hydroxymethyl-carbapenem and the 2,3-dihydro-naphtho[1,8-de][1,2]thiazine 1,1-dioxide or 1,1-dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl side chain group in a suitable solvent such as tetrahydrofuran (THF), diethyl ether, acetonitrile, dimethylformamide (DMF), benzene, dichloromethane, chloroform, and the like with a (premixed) suitable activating reagent such as diethyl azodicarboxylate (DEAD)/triphenylphosphine, diisopropyl azodicarboxylate (DIAD)/tributylphosphine, and the like, at a temperature between about -20°C and 35°C for about 5 to 90 minutes.

Alternatively, the 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl side chain group and carbapenem can be mixed together with either the azodicarboxylate or the phosphine reagent in a suitable solvent and the other component of the activating reagent (the phosphine or the azodicarboxylate, respectively) can be added to that mixture. Once the 1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl side chain group carbapenem and activating reagent(s) have been mixed, the reaction is allowed to proceed at a temperature between about -20 °C and 35 °C for about 5 to 90 minutes.

The resulting mixture is then subjected to a standard work-up procedure familiar to those skilled in the art to afford a crude 2- (1,1 dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl or 1,1,3 trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-yl) methyl substituted carbapenem

which is purified, if necessary, by recrystallization or by chromatography on silica gel, eluting with a suitable solvent or mixture of two or more solvents, such as hexane, ethyl acetate, ether, benzene, dichloromethane, chloroform, acetone, methanol and the like.

5 In some cases, it is necessary to modify substituent R* in intermediate A2 to produce the desired substituent R in final product I. Modification of the R* substituent is generally best accomplished before the removal of the protecting groups P* and R^o. This process is illustrated in Flow Chart B in which intermediate B1 is an example of 10 intermediate A2 wherein one of the R* groups is a precursor to the substituent L-Q-R^q. The skilled artisan should note that the substituent L-Q-R^q can be located in either of the phenyl rings of the final product Ib even though the scheme illustrates the process for one ring only.

15 In intermediate B1 of Flow Chart B, the substituent R* is represented as L-OP", wherein L is as previously defined and P" is a hydroxyl protecting group. Suitable protecting groups P" are trimethylsilyl and triethylsilyl. The trialkylsilyl group is easily removed by treatment with a strong acid such as trifluoromethane sulfonic acid, sulfuric acid, hydrochloric acid, or the like in a solvent consisting of 20 water plus a miscible organic solvent such as tetrahydrofuran, acetonitrile, or isopropanol. A positively charged substituent may be introduced into the side chain by first activating the hydroxyl group of L-OH by converting it to a suitable leaving group such as a triflate, mesylate, tosylate, iodide, chloride, bromide, and the like, and then 25 displacing the resulting leaving group with a compound Q'-R^{q'} such as N-methyl-imidazole, N-(2-hydroxyethyl)-imidazole, 1-methyl-4-aza-1-azoniabicyclo [2.2.2] octane, 1-(carbamoylmethyl)-4-aza-1-azoniabicyclo-[2.2.2]-octane, 1-(3-hydroxyprop-1-yl)-4-aza-1-azoniabicyclo-[2.2.2]-octane, and the like which contains a nitrogen atom that can act as a 30 nucleophile.

35 In some cases, activation of the hydroxyl group and displacement by Q'-R^{q'} to produce B3 may be accomplished in a single step by taking advantage of the basic character of compound Q'-R^{q'} and using it as a base in the activation reaction.

35 The conversion of the hydroxyl group to a suitable

leaving group is accomplished by treating the hydroxyl substituted compound in a suitable solvent such as dichloromethane, tetrahydrofuran, ether, benzene, and the like with an activating reagent, such as trifluoromethanesulfonic anhydride, methanesulfonic anhydride, 5 toluenesulfonic anhydride, methanesulfonyl chloride, benzenesulfonyl chloride, toluenesulfonyl chloride, and the like in the presence of a suitable base such as triethylamine, tributylamine, diisopropylethylamine, and the like at a temperature between about -100°C and 0°C for about 5 to 120 minutes. The intermediate thus obtained contains a 10 leaving group, which may be converted to an alternative leaving group, iodide, by treating a solution of the intermediate in a suitable solvent such as acetone, methyl ethyl ketone, and the like at about -10°C to 50°C with an excess of sodium iodide or potassium iodide for about 0.25 to 24 hours.

15 In many cases, the iodide is obtained in sufficiently pure form that it may be used without further purification. For ease of handling, the iodide, if not crystalline, may be lyophilized from benzene to afford an amorphous, easily handled, solid.

15 The activated hydroxyl group or iodide is displaced by 20 reacting the activated intermediate with reagent Q'-R^q. In some cases, activation and displacement of the hydroxyl group may be accomplished in a single step. The activating reagent is added to a solution of the hydroxyl substituted compound in the presence of a suitable base in a suitable solvent such as dichloromethane, tetrahydrofuran, ether, DMF, 25 benzene, acetonitrile, DMSO, and the like as described in the preceding paragraphs. The resulting activated intermediate is treated with 1-3 molar equivalents of compound Q'-R^q at a temperature between about -78°C and 50°C for about 15 to 120 minutes. In some cases, it is desirable to form the activated intermediate in one solvent, isolate the activated 30 intermediate, and conduct the displacement reaction in a different solvent. In other cases, the displacement may be conducted without isolation of the intermediate and, in cases where Q'-R^q is also used as a base, may even be concurrent with the formation of the activated intermediate.

In cases where the displacement reaction is best accomplished by using the iodide, a solution of the iodide is combined with an approximately equivalent amount (0.9 - 1.05 molar equivalents) of compound Q'-R^q. A silver salt of a non-nucleophilic acid, such as 5 silver trifluoromethanesulfonate, silver tetrafluoroborate and the like is then added. Although the reaction will proceed in the absence of the silver salt, the reaction proceeds more rapidly in the presence of the silver salt. In addition, the silver salt assists in the removal of the displaced iodide from the reaction mixture which can improve the 10 efficiency of subsequent steps. The resulting mixture is then subjected to a standard work-up procedure familiar to those skilled in the art to afford a crude product which is purified, if necessary, by recrystallization or chromatography.

An alternative method for introducing a positive charge into 15 the side chain may be applied to side chains (i.e. R^{*} groups) that contain a nitrogen atom which may be quaternized by reaction with a suitable alkylating reagent AR, such as methyl iodide, methyl bromide, benzyl trichloroacetimidate, methyl trifluoromethanesulfonate, triethyloxonium tetrafluoroborate, and the like. Quaternization of the 20 nitrogen atom in the side chain is effected by treating a solution of the compound with a slight excess (1.05 to 1.2 molar equivalents) of the alkylating reagent.

Modification of the substituent R^{q'}, if necessary, and removal of the remaining protecting group(s) affords the final product 25 (Ib). These transformations can be accomplished by a number of well known techniques depending on the protecting groups employed or the modifications required to transform R^{q'} to R^q or the protecting groups employed.

For example, when R^q of the final product contains a 30 primary amino group (-NH₂ or -N⁺H₃ in its protonated form), the amino group would normally be present in R^{q'} as a protected form or as a precursor group. Suitably protected forms of the amino group are allyloxycarbonylamino or p-nitrobenzyloxycarbonylamino whereas as 35 suitable amine precursor is an azido group. The amino group is liberated by established methodology either concurrent with or after the

deblocking of P^* and R^o . For example, the azido and p-nitrobenzyloxycarbonylamino groups can be converted to the amino group by catalytic hydrogenation and the allyloxycarbonylamino group provides the amino substituent on treatment with a palladium catalyst 5 in the presence of an allyl scavenger.

The synthesis of the target compound is completed by removing any protecting groups which are present in the penultimate intermediate using standard techniques which are well known to those skilled in the art. The deprotected final product is then purified, as 10 necessary, using standard techniques such as ion exchange chromatography, HPLC on reverse phase silica gel, MPLC on reverse phase polystyrene gel, and the like or by recrystallization.

The final product may be characterized structurally by standard techniques such as NMR, IR, MS, and UV. For ease of 15 handling, the final product, if not crystalline, may be lyophilized from water to afford an amorphous, easily handled solid.

The compounds of the present invention are valuable 20 antibacterial agents active against various Gram-positive and to a lesser extent Gram-negative bacteria, and accordingly find utility in human and veterinary medicine.

Many of compounds of the present invention are 25 biologically active against MRSA/MRCNS. In vitro antibacterial activity is predictive of in vivo activity when the compounds are administered to a mammal infected with a susceptible bacterial organism.

Using standard susceptibility tests, the compounds of the invention are determined to be active against MRSA.

The compounds of the invention can be formulated in 30 pharmaceutical compositions by combining the compound with a pharmaceutically acceptable carrier. Examples of such carriers are set forth below.

The compounds may be employed in powder or crystalline form, in liquid solution, or in suspension. They may be administered by a variety of means; those of principal interest

include: topically, orally and parenterally by injection (intravenously or intramuscularly).

Compositions for injection, a preferred route of delivery, may be prepared in unit dosage form in ampules, or in multidose 5 containers. The injectable compositions may take such forms as suspensions, solutions, or emulsions in oily or aqueous vehicles, and may contain various formulating agents. Alternatively, the active ingredient may be in powder (lyophilized or non-lyophilized) form for reconstitution at the time of delivery with a suitable vehicle, such 10 as sterile water. In injectable compositions, the carrier is typically comprised of sterile water, saline or another injectable liquid, e.g., peanut oil for intramuscular injections. Also, various buffering agents, preservatives and the like can be included.

Topical applications may be formulated in carriers such 15 as hydrophobic or hydrophilic bases to form ointments, creams, lotions, in aqueous, oleaginous or alcoholic liquids to form paints or in dry diluents to form powders.

Oral compositions may take such forms as tablets, capsules, oral suspensions and oral solutions. The oral compositions 20 may utilize carriers such as conventional formulating agents, and may include sustained release properties as well as rapid delivery forms.

The dosage to be administered depends to a large extent upon the condition and size of the subject being treated, the route and 25 frequency of administration, the sensitivity of the pathogen to the particular compound selected, the virulence of the infection and other factors. Such matters, however, are left to the routine discretion of the physician according to principles of treatment well known in the antibacterial arts. Another factor influencing the precise dosage 30 regimen, apart from the nature of the infection and peculiar identity of the individual being treated, is the molecular weight of the compound.

The compositions for human delivery per unit dosage, whether liquid or solid, may contain from about 0.01% to as high as 35 about 99% of active material, the preferred range being from about 10-

60%. The composition will generally contain from about 15 mg to about 2.5 g of the active ingredient; however, in general, it is preferable to employ dosage amounts in the range of from about 250 mg to 1000 mg. In parenteral administration, the unit dosage will 5 typically include the pure compound in sterile water solution or in the form of a soluble powder intended for solution, which can be adjusted to neutral pH and isotonic.

The invention described herein also includes a method of treating a bacterial infection in a mammal in need of such treatment 10 comprising administering to said mammal a compound of formula I in an amount effective to treat said infection.

The preferred methods of administration of the Formula I antibacterial compounds include oral and parenteral, e.g., i.v. infusion, i.v. bolus and i.m. injection.

15 For adults, about 5-50 mg of Formula I antibacterial compound per kg of body weight given one to four times daily is preferred. The preferred dosage is 250 mg to 1000 mg of the antibacterial given one to four times per day. More specifically, for mild infections a dose of about 250 mg two or three times daily is 20 recommended. For moderate infections against highly susceptible gram positive organisms a dose of about 500 mg three or four times daily is recommended. For severe, life-threatening infections against organisms at the upper limits of sensitivity to the antibiotic, a dose of about 1000-2000 mg three to four times daily may be recommended.

25 For children, a dose of about 5-25 mg/kg of body weight given 2, 3, or 4 times per day is preferred; a dose of 10 mg/kg is typically recommended.

The compounds of Formula I are of the broad class known as carbapenems. Many carbapenems are susceptible to attack 30 by a renal enzyme known as dehydropeptidase (DHP). This attack or degradation may reduce the efficacy of the carbapenem antibacterial agent. Many of the compounds of the present invention, on the other hand, are less subject to such attack, and therefore may not require the use of a DHP inhibitor. However, such use is optional and 35 contemplated to be part of the present invention. Inhibitors of DHP

and their use with carbapenems are disclosed in, e.g., [European Patent Application Nos. 79102616.4, filed July 24, 1979 (Patent No. 0 007 614); and 82107174.3, filed August 9, 1982 (Publication No. 0 072 014)].

5 The compounds of the present invention may, where DHP inhibition is desired or necessary, be combined or used with the appropriate DHP inhibitor as described in the aforesaid patents and published application. The cited European Patent Applications define the procedure for determining DHP susceptibility of the
 10 present carbapenems and disclose suitable inhibitors, combination compositions and methods of treatment. A preferred weight ratio of Formula I compound: DHP inhibitor in the combination compositions is about 1:1.

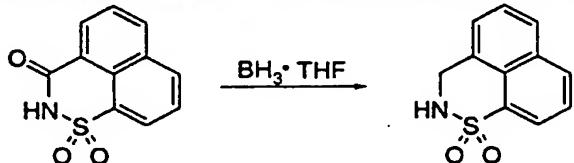
15 A preferred DHP inhibitor is 7-(L-2-amino-2-carboxy-ethylthio)-2-(2,2-dimethylcyclopropanecarboxamide)-2-heptenoic acid or a useful salt thereof.

The invention is further described in connection with the following non-limiting examples.

20

PREPARATIVE EXAMPLE 1

2,3-DIHYDRO-NAPHTHO[1,8-DE][1,2]THIAZINE 1,1-DIOXIDE



25 A solution of 1,1-dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-3-one (137 mg, 0.587 mmol) in anhydrous tetrahydrofuran (2 mL) was treated with a 1.0M solution of borane-tetrahydrofuran complex in tetrahydrofuran (1.17 mL, 1.17 mmol). The resulting solution was heated at 65°C for 5 hours. The reaction mixture was quenched with methanol (5 mL) and evaporated under vacuum. The
 30 residue was purified by preparative layer chromatography on two 1mm

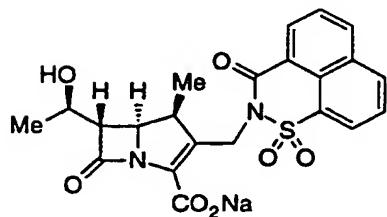
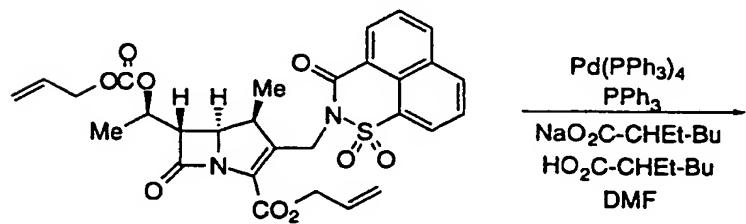
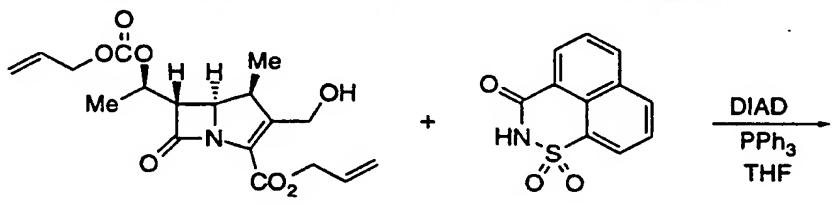
x 20cm x 20cm silica gel GF plates, developing with 7:3 ethyl acetate-hexanes, to afford the title compound (78 mg, 61% yield).

The 1,1-dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-3-one starting material can be prepared according to the procedure described in Lombardino, J. G. *J. Org. Chem.* 1971, 36, 1843.

EXAMPLE 1

SODIUM (1S,5R,6S)-6-[1(R)-HYDROXY-ETHYL]-1-METHYL-2-(1,1,3-TRIOXO-2,3-DIHYDRO-NAPHTHO[1,8-DE][1,2]THIAZIN-2-

YLMETHYL)-CARBAPEN-2-EM-3-CARBOXYLATE



Step 1: Allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-1-methyl-2-(1,1,3-trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl)-carbapen-2-em-3-carboxylate

A solution of allyl (1*S*,5*R*,6*S*)-6-[1(*R*)-allyloxycarbonyloxy-thyl]-2-hydroxymethyl-1-methyl-carbapen-2-em-3-carboxylate (64.8 mg, 0.174 mmol), 1,1-dioxo-2,3-dihydro-naphtho[1,8-*de*][1,2]thiazin-3-one (34.5 mg, 0.145 mmol), and triphenylphosphine (46 mg, 0.174 mmol) in
5 anhydrous tetrahydrofuran (2 mL) was cooled in an ice bath and stirred under a nitrogen atmosphere. Diisopropyl azodicarboxylate (0.034 mL, 0.174 mmol) was added and the resulting mixture was stirred at 0-5°C for 10 minutes. The mixture was concentrated under vacuum and the residue was purified by preparative layer chromatography on a 1mm x
10 20cm x 20 cm silica gel GF plate using 7:3 ethyl acetate-hexanes as the developing solvent. The product band was removed and eluted with ethyl acetate to afford allyl (1*S*,5*R*,6*S*)-6-[1(*R*)-allyloxycarbonyloxy-ethyl]-1-methyl-2-(1,1,3-trioxo-2,3-dihydro-naphtho[1,8-*de*][1,2]thiazin-2-ylmethyl)-carbapen-2-em-3-carboxylate (49 mg) as an oil.

15

Step 2: Sodium (1*S*,5*R*,6*S*)-6-[1(*R*)-hydroxy-ethyl]-1-methyl-2-(1,1,3-trioxo-2,3-dihydro-naphtho[1,8-*de*][1,2]thiazin-2-ylmethyl)-carbapen-2-em-3-carboxylate

The product from step 1 (56 mg, 0.0961 mmol),
20 triphenylphosphine (15 mg, 0.0577 mmol), 0.5M sodium 2-ethyl-hexanoate in ethyl acetate (0.211 mL, 0.106 mmol), 2-ethyl-hexanoic acid (0.0146 mL, 0.106 mmol), and tetrakis(triphenylphosphine)palladium(0) (22 mg, 0.0192 mmol) were combined in anhydrous dimethylformamide (1 mL). The mixture was stirred at room temperature for 5 minutes,
25 then cooled in an ice bath and stirred at 0-5°C for 2 hours. The mixture was diluted with diethyl ether (12 mL) to precipitate a solid. The solid was collected by centrifugation, washed with ether (5 mL), and dried under vacuum. The solid was dissolved in water (5 mL) and added to a column (1 x 20 cm) of Amberchrom CG-161 resin. The column was
30 eluted with water followed by a progressive gradient of acetonitrile in water. The product containing fractions were combined and lyophilized

to a white, amorphous solid (18 mg). This material was further purified by preparative layer chromatography on a 1mm x 20cm x 20 cm RPSF plate, developing with 7:3 water-acetonitrile in an ice cooled tank. The product band was located by UV, removed and eluted with 4:1

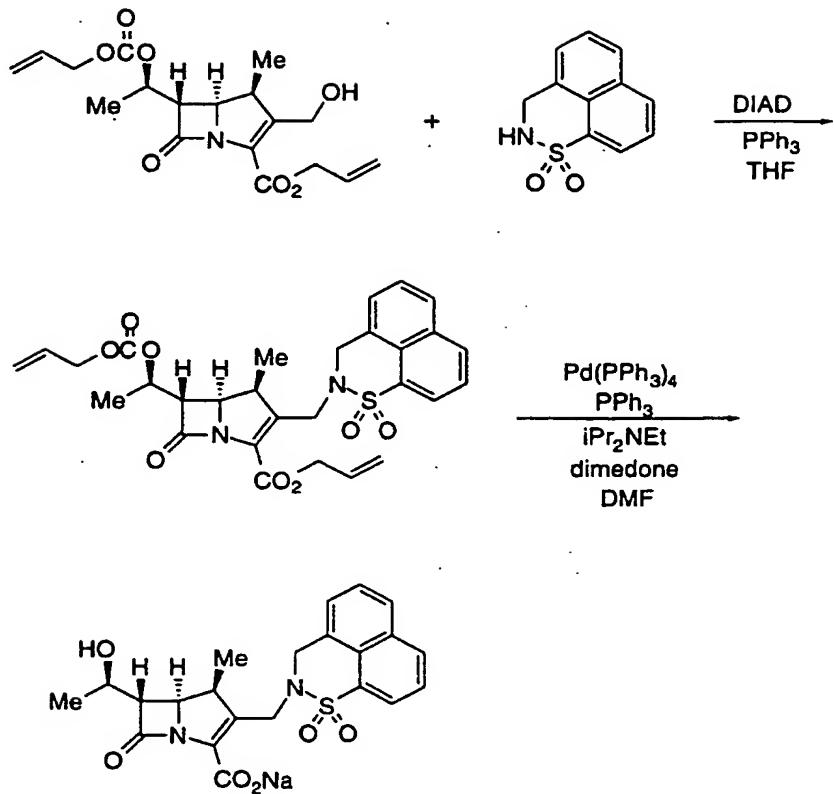
5 acetonitrile-water. The eluent was washed with hexanes, filtered through a 0.45 micron disc, concentrated under vacuum to remove the acetonitrile, and lyophilized to afford the title compound (17.9 mg) as a white amorphous solid.

10 ^1H NMR (D_2O , 500 Mhz) δ 1.33 (d, 1- CH_3), 1.36 (d, CH_3CHOH), 3.32 (m, H-1), 3.52 (dd, H-6), 4.27 (dd, H-5), 4.32 (m, CH_3CHOH), 4.82 and 5.60 (two d's, 2- CH_2), 8.03 (m, two aryl-H), 8.11 (d, aryl-H), 8.33 (d, aryl-H), 8.38 (d, aryl-H), and 8.56 (d, aryl-H)

15

EXAMPLE 2

SODIUM (1S,5R,6S)-2-(1,1-DIOXO-2,3-DIHYDRO-NAPHTHO[1,8-DE][1,2]THIAZIN-2-YLMETHYL)-6-[1(R)-HYDROXY-ETHYL]-1-METHYL-CARBAPEN-2-EM-3-CARBOXYLATE



Step 1: Allyl (1*S*,5*R*,6*S*)-6-[1(*R*)-allyloxycarbonyloxy-ethyl]-2-(1,1-dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl)-1-methyl-carbapen-2-em-3-carboxylate

5

A solution of allyl (1*S*,5*R*,6*S*)-6-[1(*R*)-allyloxycarbonyloxy-ethyl]-2-hydroxymethyl-1-methyl-carbapen-2-em-3-carboxylate (116 mg, 0.317 mmol), 2,3-dihydro-naphtho[1,8-de][1,2]thiazine 1,1-dioxide (58 mg, 0.265 mmol), and triphenylphosphine (83 mg, 0.317 mmol) in anhydrous tetrahydrofuran (4 mL) was cooled in an ice bath and stirred under a nitrogen atmosphere. Diisopropyl azodicarboxylate (0.062 mL, 0.317 mmol) was added dropwise over one minute. The resulting mixture was stirred at 0-5°C for 30 minutes. The mixture was concentrated under vacuum and the residue was purified by preparative layer

chromatography on two 1mm x 20cm x 20 cm silica gel GF plate using 1:1 ethyl acetate-hexanes as the developing solvent. The product band was removed and eluted with ethyl acetate to afford a mixture (160 mg) of allyl (1*S*,5*R*,6*S*)-6-[1(*R*)-allyloxycarbonyloxy-ethyl]-2-(1,1-dioxo-2,3-dihydro-naphtho[1,8-*de*][1,2]thiazin-2-ylmethyl)-1-methyl-carbapen-2-em-3-carboxylate and diisopropyl hydrazodiester.

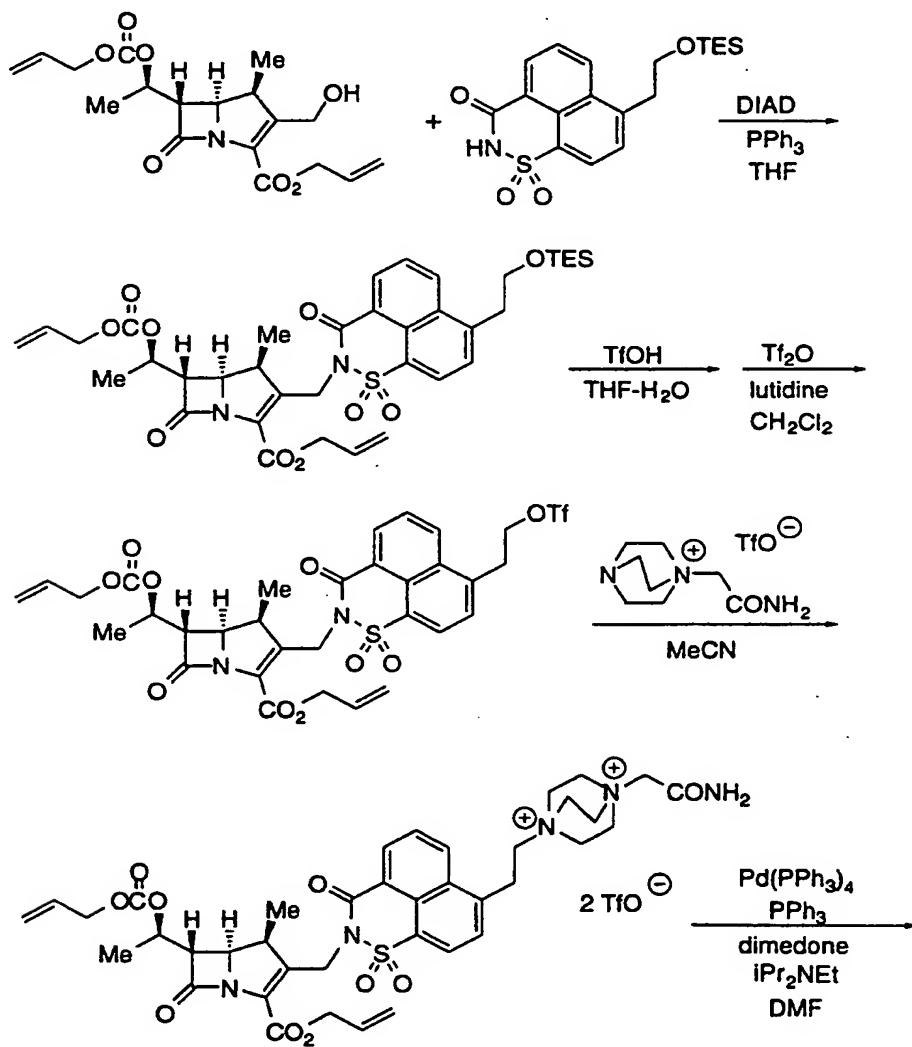
5 **Step 2: Sodium (1*S*,5*R*,6*S*)-2-(1,1-dioxo-2,3-dihydro-naphtho[1,8-*de*][1,2]thiazin-2-ylmethyl)-6-[1(*R*)-hydroxy-ethyl]-1-methyl-carbapen-2-em-3-carboxylate**

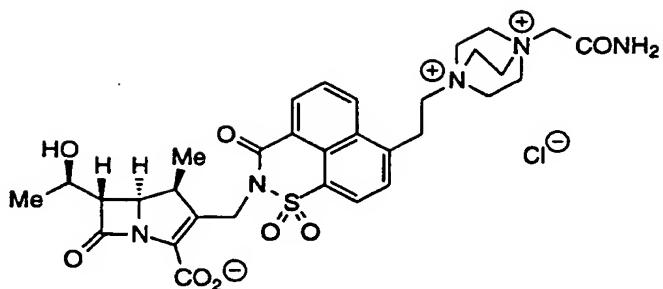
10 The product from step 1 (160 mg, ca. 0.265 mmol carbapenem component), triphenylphosphine (39 mg, 0.0398 mmol), dimedone (111 mg, 0.795 mmol), tetrakis(triphenylphosphine)-15 palladium(0) (15 mg, 0.0133 mmol) and 1.0N aqueous sodium bicarbonate (0.265 mL, 0.265 mmol) were combined in dimethylformamide (1 mL). The mixture was stirred while N,N-diisopropylethylamine (0.138 mL, 0.795 mmol) was added dropwise over one minute. After stirring for 20 minutes at room temperature, the 20 mixture was diluted with diethyl ether (10 mL) to precipitate a solid. The solid was collected by centrifugation and dried under vacuum. The solid was purified by preparative layer chromatography on a 1mm x 20cm x 20 cm RPSF plate, developing with 4:1 water-acetonitrile. The product band was located by UV, removed and eluted with 4:1 acetonitrile-water. The 25 eluent was washed with hexanes, filtered through a 0.45 micron disc, concentrated under vacuum to remove the acetonitrile, and lyophilized to afford the title compound (26 mg) as a white amorphous solid.

30 ¹H NMR (2:1 D₂O-CDCl₃, 500 MHz) δ 1.43 (d, 1-CH₃), 1.48 (d, CH₃CHOH), 3.51 (m, H-1), 3.62 (dd, H-6), 4.45 (m, H-5, CH₃CHOH and CHH), 4.75 (d, CHH), 4.83 (d, CHH), 5.04 (d, CHH), and 7.8-8.55 (six aryl-H)

EXAMPLE 3

(1S,5R,6S)-2-[7-[2-(4-CARBAMOYLMETHYL-1,4-DIAZONIA-
BICYCLO[2.2.2]OCT-1-YL)-ETHYL]-1,1,3-TRIOXO-2,3-DIHYDRO-
5 NAPHTHO[1,8-DE][1,2]THIAZIN-2-YLMETHYL]-6-[1(R)-HYDROXY-
ETHYL]-1-METHYL-CARBAPEN-2-EM-3-CARBOXYLATE CHLORIDE





Step 1: Allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-1-methyl-2-[1,1,3-trioxo-7-(2-triethylsilyloxy-ethyl)-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl]- carbapen-2-em-3-carboxylate

5

A solution of allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-2-hydroxymethyl-1-methyl-carbapen-2-em-3-carboxylate (365 mg, 1.0 mmol), triphenylphosphine (315 mg, 1.2 mmol), and 7-(2-triethylsilyloxy-ethyl)-1,1-dioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-3-one (431 mg, 1.1 mmol) in anhydrous tetrahydrofuran (7 mL) is cooled in an ice-bath and stirred under a nitrogen atmosphere while diisopropyl azodicarboxylate (0.24 mL, 1.2 mmol) is added dropwise over a few minutes. The resulting solution is stirred in the cold for 30 minutes, then diluted with chloroform, washed with 5% aqueous sodium bicarbonate and brine, dried over sodium sulfate, filtered and evaporated under vacuum. The residue is purified by silica gel flash chromatography, eluting with hexane-ethyl acetate, to afford allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-1-methyl-2-[1,1,3-trioxo-7-(2-triethylsilyloxy-ethyl)-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl]- carbapen-2-em-3-carboxylate.

Step 2: Allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-1-methyl-2-[1,1,3-trioxo-7-[2-(trifluoro-methanesulfonyloxy)-ethyl]-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl]-carbapen-2-em-3-carboxylate

The compound from step 1 (369 mg, 0.5 mmol) in tetrahydrofuran (4.0 mL) is diluted with water (1.0 mL and treated with 1M aqueous trifluoromethanesulfonic acid (0.05 mL, 0.05 mmol). After 5 stirring at room temperature for 15 minutes, the reaction mixture is partitioned between ethyl acetate (25 mL) and 5% aqueous sodium bicarbonate (5 mL). The organic phase is washed with 50% saturated brine, dried over magnesium sulfate, filtered, evaporated under vacuum, and stripped with anhydrous toluene to leave a residue of the 10 desilylated alcohol.

A solution of the crude alcohol (0.5 mmol) in anhydrous dichloromethane (10 mL) is cooled in an ice-methanol bath (-20°C) and stirred under a nitrogen atmosphere. The solution is treated with 2,6-lutidine (0.175 mL, 1.5 mmol) followed by trifluoromethanesulfonic 15 anhydride (0.126 mL, 0.75 mmol). After stirring at -20° to -15°C (bath temperature) for 40 minutes, the solution is diluted with dichloromethane (30 mL), washed with water (20 mL), 0.1N hydrochloric acid (20 mL), and water (20 mL), dried over magnesium sulfate, filtered, and evaporated under vacuum to afford allyl (1S,5R,6S)-6-[1(R)- 20 allyloxycarbonyloxy-ethyl]-1-methyl-2-[1,1,3-trioxo-7-[2-(trifluoromethanesulfonyloxy)-ethyl]-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl]-carbapen-2-em-3-carboxylate.

Step 3: Allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-2-[7-[2-(4-carbamoylmethyl-1,4-diazonia-bicyclo[2.2.2]oct-1-yl)-ethyl]-1,1,3-trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl]-1-methyl-carbapen-2-em-3-carboxylate bis(trifluoromethanesulfonate)

30 A sample (92 mg, 0.125 mmol) of the triflate derivative prepared as described in step 2 is dissolved in anhydrous acetonitrile (1.0 mL) and the solution is treated with 1-carbamoylmethyl-1-azonia-4-aza-

bicyclo[2.2.2]octane trifluoromethanesulfonate (44 mg, 0.14 mmol). The reaction mixture is stirred at room temperature for 90 minutes, then evaporated under vacuum to a residue which is aged an additional 90 minutes at room temperature. The residue is triturated with anhydrous diethyl ether to afford allyl (1S,5R,6S)-6-[1(R)-allyloxycarbonyloxy-ethyl]-2-[7-[2-(4-carbamoylmethyl-1,4-diazonia-bicyclo[2.2.2]oct-1-yl)-ethyl]-1,1,3-trioxo-2,3-dihydro-naphtho[1,8-de][1,2]thiazin-2-ylmethyl]-1-methyl-carbapen-2-em-3-carboxylate bis(trifluoromethanesulfonate).

10 Step 4: (1S,5R,6S)-2-[7-[2-(4-carbamoylmethyl-1,4-diazonia-
bicyclo[2.2.2]oct-1-yl)-ethyl]-1,1,3-trioxo-2,3-dihydro-naphtho[1,8-
de][1,2]thiazin-2-ylmethyl]-6-[1(R)-hydroxy-ethyl]-1-methyl-carbapen-2-
em-3-carboxylate chloride

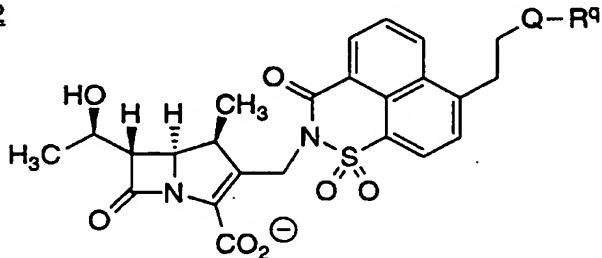
15 The crude bisprotected intermediate of step 3 (approximately 0.125 mmol), triphenylphosphine (4.9 mg, 0.0187 mmol), dimedone (53 mg, 0.378 mmol), and tetrakis(triphenylphosphine)palladium(0) (7.2 mg, 0.0062 mmol) are dissolved in anhydrous dimethylformamide (1.3 mL). The solution is purged with nitrogen, then treated with *N,N*-diisopropylethylamine (0.065 mL, 0.373 mmol). After stirring at room temperature for 15 minutes, the reaction mixture is added to diethyl ether to precipitate the crude product. The precipitate is triturated with ether (2 x 5 mL) and dried under vacuum. The precipitate in 1:1 acetonitrile-water (1 mL) is added to a column of Macro-Prep CM (Bio-Rad) weak cation exchange resin (3 mL). The column is successively eluted with 1:1 acetonitrile-water (4 mL), water (3 x 5 mL), and 5% aqueous sodium chloride (5 x 3 mL). The product containing sodium chloride fractions are pooled, cooled in ice, and loaded onto a column of Amberchrom CG-161 (TosoHaas) resin (3 mL). The Amberchrom column is eluted with ice-cold water (3 x 5 mL) followed by 20% isopropanol in water (10 x 3 mL).

The product containing aqueous isopropanol fractions are combined, concentrated under vacuum to remove the isopropanol, and lyophilized to afford the title compound as an amorphous solid.

EXAMPLES 4-32

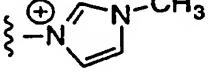
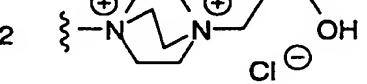
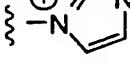
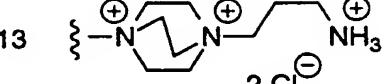
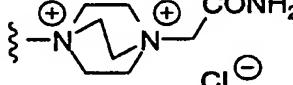
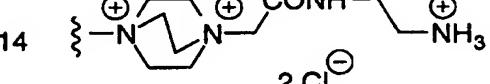
By appropriately modifying the procedures of Examples 1-3, the following compounds are prepared:

TABLE 2



Ex.	Q-R ^q	Ex.	Q-R ^q
4	$\text{---N}^+ \text{---CH}_3$	7	$\text{---N}^+ \text{---CH}_2 \text{---CH}_2 \text{---NH}_3^+$ 2 Cl ⁻
5	$\text{---N}^+ \text{---CH}_2 \text{---CH}_2 \text{OH}$	8	$\text{---N}^+ \text{---CH}_2 \text{---CONH---CH}_2 \text{---NH}_3^+$ 2 Cl ⁻
6	$\text{---N}^+ \text{---CH}_2 \text{---CH}_2 \text{---OH}$ Cl ⁻		

TABLE 3

Ex.	Q-R ^q	Ex.	Q-R ^q
9		12	
10		13	
11		14	

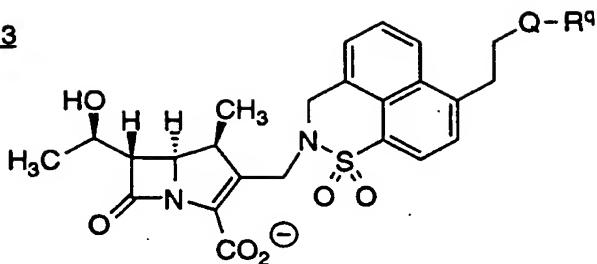
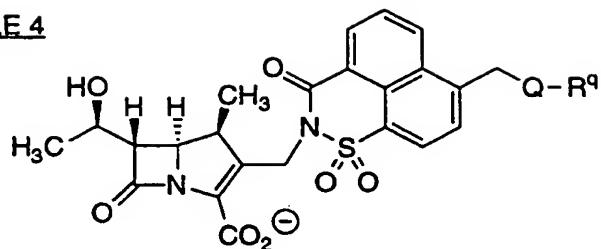
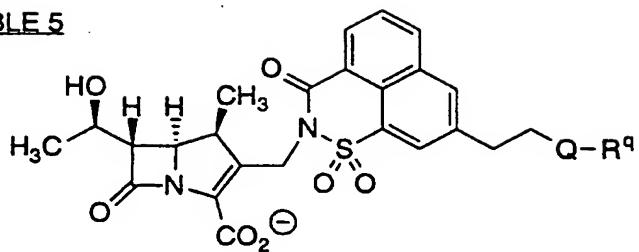


TABLE 4



Ex.	Q-R ^q	Ex.	Q-R ^q
15		18	
16		19	
17		20	

TABLE 5



Ex.

Q-R^q

Ex.

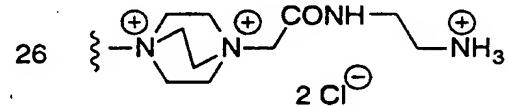
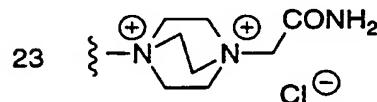
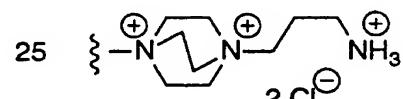
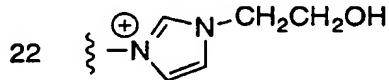
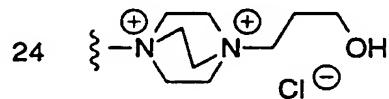
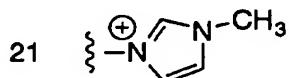
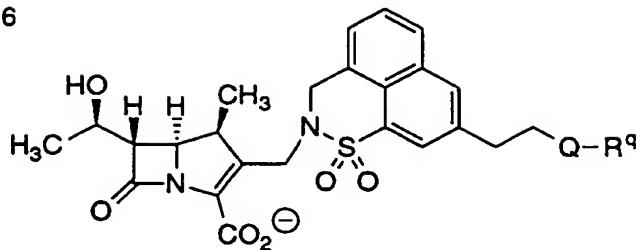
Q-R^q

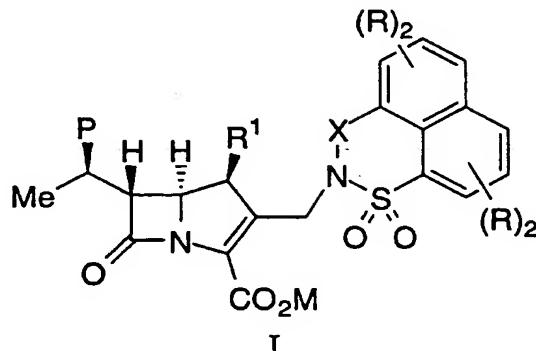
TABLE 6



Ex.	Q-R ^q	Ex.	Q-R ^q
27		30	
28		31	
29		32	

WHAT IS CLAIMED IS:

1. A compound represented by formula I:



including pharmaceutically acceptable salts thereof, wherein:

R¹ represents H or methyl;

10

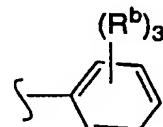
CO₂M represents a carboxylic acid, a pharmaceutically acceptable carboxylic acid salt, carboxylate anion, a pharmaceutically acceptable ester group or a carboxylic acid protected by a protecting group;

15

X represents CH₂ or C=O;

P represents hydrogen, hydroxyl, F or hydroxyl protected by a hydroxyl-protecting group;

20

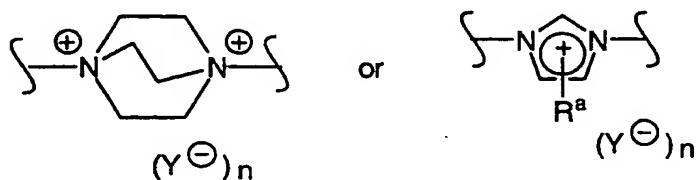
each R independently represents R^b,
 -Het(R^b)₃,
 C₂₋₆ alkenyl, or a group L-Q-R^a with the proviso that only one R group of the type L-Q-R^a can be present or one R group may be taken with L, if present, and any intervening atoms to represent a 5-6 membered ring;

25

L is C₁₋₄ straight or branched alkylene, uninterrupted, interrupted or terminated by 1-2 of O, S, NRA, C(O), CO₂ and C(O)NRA;

Q represents:

5



Y- is a charge balancing group;

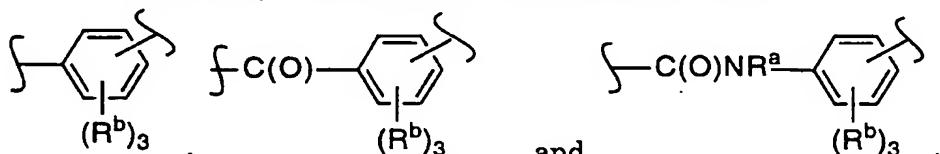
n is a value from 0 to 2, selected to maintain overall charge

10 neutrality;

R^a is H or C₁₋₆ alkyl;

R^q is C₁₋₆ alkyl, straight or branched, uninterrupted, interrupted or terminated by 1-2 of O, S, NRA, C(O), C(O)O, C(O)NRA^a, -CH=CH-, -Het(R^b)₃-, -C(O)Het(R^b)₃-, -C(O)NRA^aHet(R^b)₃-,

15



said R^q being unsubstituted or substituted with 1-3 R^c groups;

20

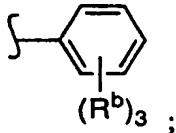
Het is a heteroaryl group;

each R^b is independently selected from H, halo, OR^a, OC(O)R^a, C(O)R^a, CN, C(O)NRA^aRd, NO₂, NRA^aRd, SO₂NRA^aRd and C₁₋₄ alkyl unsubstituted or substituted with 1-3 groups selected from R^e;

25

each R^c is independently selected from halo, OR^f, OC(O)R^f, SR^f, S(O)R^f, SO₂R^f, CN, C(O)R^f, CO₂R^f, NR^fRg, N⁺R^aR^fR^fZ⁻, C(O)NRA^aR^f, -Het(R^b)₃, C(=N⁺R^aR^f)RaZ⁻, C(=N⁺R^aR^f)NRA^aR^fZ⁻,

NR_aC(=N⁺R_aR_f)R_aZ⁻, NR_aC(=N⁺R_aR_f)NR_aR_fZ⁻, heteroarylium(R_b)₃Z⁻, SO₂NR_aR_f, OC(O)NR_aR_f, NR_aC(O)R_f, NR_aC(O)NR_aR_f, and

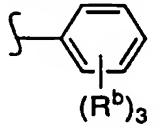


5 or in the alternative, when 2 or more R^c groups are present, 2 R^c groups may be taken together with any intervening atoms to form a 3-6 membered carbocyclic ring, optionally interrupted with 1-3 of O, S, NR_g, and C(O), said ring being unsubstituted or substituted with 1-3 R^e groups;

10 R^d is H or C₁₋₄ alkyl, or R^a and R^d taken together with any intervening atoms represent a 4-6 membered ring;

15 R^e is selected from halo, OR_a, NR_aR^d and CONR_aR^d;

15 R^f is H; C₁₋₆ straight or branched chain alkyl, unsubstituted or substituted with 1-3 R^e groups; -Het(R_b)₃; C₃₋₆ cycloalkyl, unsubstituted or substituted with 1-3 R^e groups, and



20 or R^a and R^f together with an intervening atoms form a 4-6 membered ring, optionally interrupted by O, S, NR_a or C(O); and

25 R^g is H, C₁₋₆ alkyl, unsubstituted or substituted with 1-3 R^e groups; C₃₋₆ cycloalkyl, unsubstituted or substituted with 1-3 R^e groups; C(=N⁺R_aR_f)R_aZ⁻ or C(=N⁺R_aR_f)NR_aR_fZ⁻;

or R^f and R^g together with any intervening atoms form a 4-6 membered ring optionally interrupted by O, S, NR_a or C(O); and

Z^- is a charge balancing group selected from Y^- , a monovalent anion such as acetate, benzoate, bromide, chloride or the like, or an internal anion such as CO_2M , where M is a negative charge.

5

2. A compound in accordance with claim 1
wherein R^1 represents methyl.

10 3. A compound in accordance with claim 1
wherein CO_2M represents a carboxylate anion.

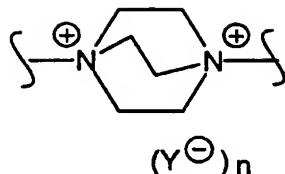
15 4. A compound in accordance with claim 1
wherein one R represents $L-Q-Rq$ and all other R groups represent H,
halo or C_{1-4} alkyl unsubstituted or substituted with 1-3 groups selected
from R_e .

20 5. A compound in accordance with claim 4
wherein M represents a negative charge which is balanced by a
positively charged Q group.

25 6. A compound in accordance with claim 4
wherein if the positively charged Q group contains more than one
positive charge, a negatively charged counterion may be present which
in combination with the carboxylate anion, provides overall charge
neutrality.

30 7. A compound in accordance with claim 4
wherein L represents $-CH_2-$ or $-CH_2CH_2-$.

8. A compound in accordance with claim 4
wherein Q represents



wherein Y- represents a charge balancing group, and n is 0 to 2.

9. A compound in accordance with claim 4
 5 wherein Rq is straight or branched C₁₋₆ alkyl, substituted with 1-3 Rc groups.

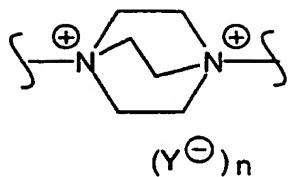
10. A compound in accordance with claim 1
 wherein X is CH₂.

11. A compound in accordance with claim 1
 wherein X is CO.

12. A compound in accordance with claim 1
 15 wherein each R is independently H, halo or C₁₋₄ alkyl unsubstituted or substituted with 1-3 groups selected from Re.

13. A compound in accordance with claim 1
 wherein:
 20 R¹ represents CH₃;
 CO₂M represents a carboxylate anion;
 P represents hydroxyl or hydroxyl protected by a hydroxyl protecting group;
 one R is L-Q-R^a and each remaining R is independently H, halo or C₁₋₄ alkyl unsubstituted or substituted with 1-3 groups selected from Re;
 25 R^a is H or C₁₋₆ alkyl;
 Rd is H or C₁₋₄ alkyl, or R^a and Rd taken together with any intervening atoms represent a 4-6 membered ring;
 30 Re is halo, OR^a, NR^aRd or CONR^aRd;
 L represents -CH₂- or -CH₂CH₂-;

Q represents



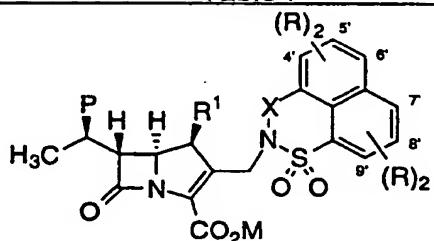
wherein Y- represents a charge balancing group, and n is 0 to 2; and

5 R₉ is straight or branched C₁₋₆ alkyl, optionally interrupted

by $C(O)NR^a$ or  and
substituted with 1-3 R^c groups, and
 R^c is as originally defined.

10 14. A compound in accordance with claim 1
represented by the formulas in Table 1:

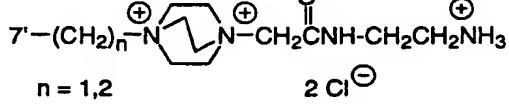
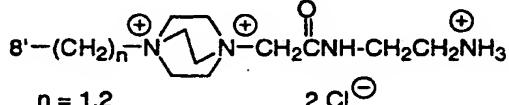
Table 1



P = H, F or OH, and R¹ = Methyl or H

	X	Substituent(s) R	M
1	CO	H	Na
2	CH ₂	H	Na
3	CO	7'-(CH ₂) _n -OH	Na
4	CO	7'-(CH ₂) _n - N^{\oplus}  -CH ₃ n = 1,2	⊖
5	CO	8'-(CH ₂) _n - N^{\oplus}  -CH ₃ n = 1,2	⊖
6	CH ₂	7'-(CH ₂) _n - N^{\oplus}  -CH ₃ n = 1,2	⊖
7	CO	7'-(CH ₂) _n - N^{\oplus}  -CH ₂ CH ₂ OH n = 1,2	⊖

	X	Substituent(s) R	M
16	CO	$7'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{OH}$ $n = 1, 2$	\ominus
17	CO	$8'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{OH}$ $n = 1, 2$	\ominus
18	CH_2	$7'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{OH}$ $n = 1, 2$	\ominus
19	CO	$7'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{NH}_3$ $n = 1, 2$ 2Cl^-	\ominus
20	CO	$8'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{NH}_3$ $n = 1, 2$ 2Cl^-	\ominus
21	CH_2	$7'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{NH}_3$ $n = 1, 2$ 2Cl^-	\ominus
22	CO	$7'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}\text{CH}_2\text{---}\overset{\text{O}}{\text{C}}\text{---}\text{CNH---CH}_2\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{NH}_3$ $n = 1, 2$ 2Cl^-	\ominus
23	CO	$8'-(\text{CH}_2)_n-\overset{+}{\text{N}}\text{---}\text{C}_6\text{H}_4\text{---}\overset{+}{\text{N}}\text{---}\text{CH}_2\text{---}\overset{\text{O}}{\text{C}}\text{---}\text{CNH---CH}_2\text{---}\overset{+}{\text{N}}\text{---}(\text{CH}_2)_3\text{NH}_3$ $n = 1, 2$ 2Cl^-	\ominus

	X	Substituent(s) R	M
24	CH ₂		⊖
25	CH ₂		⊖
26	CO	6'-Br, 8'-(CH ₂) _n -N+(cyclohexyl)-N+(CH ₂)CONH ₂ Cl ⁻ where n=1,2	⊖
27	CO	6'-Br, 8'-(CH ₂) _n -N+(cyclohexyl)-N+(CH ₂)CONH ₂ Cl ⁻ where n=1,2	⊖
28	CH ₂	6'-Br, 8'-(CH ₂) _n -N+(cyclohexyl)-N+(CH ₂)CONH ₂ Cl ⁻ where n=1,2	⊖
29	CO	7'-(CH ₂) _n -N+(cyclohexyl)-N+(CH ₂)CONH ₂ Cl ⁻ where n=1,2	⊖
30	CH ₂	7'-(CH ₂) _n -N+(cyclohexyl)-N+(CH ₂)CONH ₂ Cl ⁻ where n=1,2	⊖
31	CO	8'-(CH ₂) _n -N+(cyclohexyl)-N+(CH ₂)CONH ₂ Cl ⁻ where n=1,2	⊖

	X	Substituent(s) R	M
32	CH ₂	$7'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{OH})$ $n = 1, 2$ Cl^-	\ominus
33	CO	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{OH})$ $n = 1, 2$ Cl^-	\ominus
34	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{OH})$ $n = 1, 2$ Cl^-	\ominus
35	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_3)$ $n = 1, 2$ Cl^-	\ominus
36	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{OH})$ $n = 1, 2$ Cl^-	\ominus
37	CH ₂	$8'-(\text{CH}_2)_n\text{N}^+(\text{C}_6\text{H}_4\text{CH}_2\text{CH}_2\text{NH}_3^+)$ $n = 1, 2$ 2Cl^-	\ominus

15. A pharmaceutical composition comprised of a
 5 compound in accordance with claim 1 in combination with a
 pharmaceutically acceptable carrier.

16. A method of treating a bacterial infection
 comprising administering to a mammalian patient in need of such

treatment a compound as defined in claim 1 in an amount which is effective for treating a bacterial infection.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US99/14235

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C07D 477/14

US CL :540/302; 514/210

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 540/302; 514/210

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS Online

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,756,725 A (WILKENING et al.) 26 May 1998, Abstract.	1-16

Further documents are listed in the continuation of Box C.

See patent family annex.

•	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"B"	earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"A"	document member of the same patent family
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

15 SEPTEMBER 1999

Date of mailing of the international search report

21 OCT 1999

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

D. Lawrence Far
MARK L. BERCH

Telephone No. (703) 308-1235